

United States Patent [19]

Meister et al.

[11] Patent Number: **4,665,082**

[45] Date of Patent: **May 12, 1987**

[54] **CYSTEINE DELIVERY SYSTEM**

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[21] Appl. No.: **750,043**

[22] Filed: **Jun. 27, 1985**

Related U.S. Application Data

[63] Continuation of Ser. No. 547,459, Nov. 1, 1983, abandoned.

[51] Int. Cl.⁴ **A61K 31/425**; A61K 31/23

[52] U.S. Cl. **514/365**; 514/369;
514/562

[58] Field of Search 514/365, 369, 562

[56] **References Cited**

U.S. PATENT DOCUMENTS

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Kaneko et al., Bull. Chem. Soc., 37: 242-244, 1964.
PDR, 34th ed, 1980, pp. 1238-1241 and 1730-1732.

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Attorney, Agent, or Firm—Kenyon & Kenyon

[57] **ABSTRACT**

L-2-oxothiazolidine-4-carboxylate, a sulfur analog of 5-oxoproline, is cleaved by the enzyme 5-oxo-L-prolinase to form cysteine, thus providing the basis for a cysteine delivery system by the addition of L-2-oxothiazolidine-4-carboxylate to base amino acid solutions or by injecting it directly into in vivo cells.

7 Claims, 4 Drawing Figures

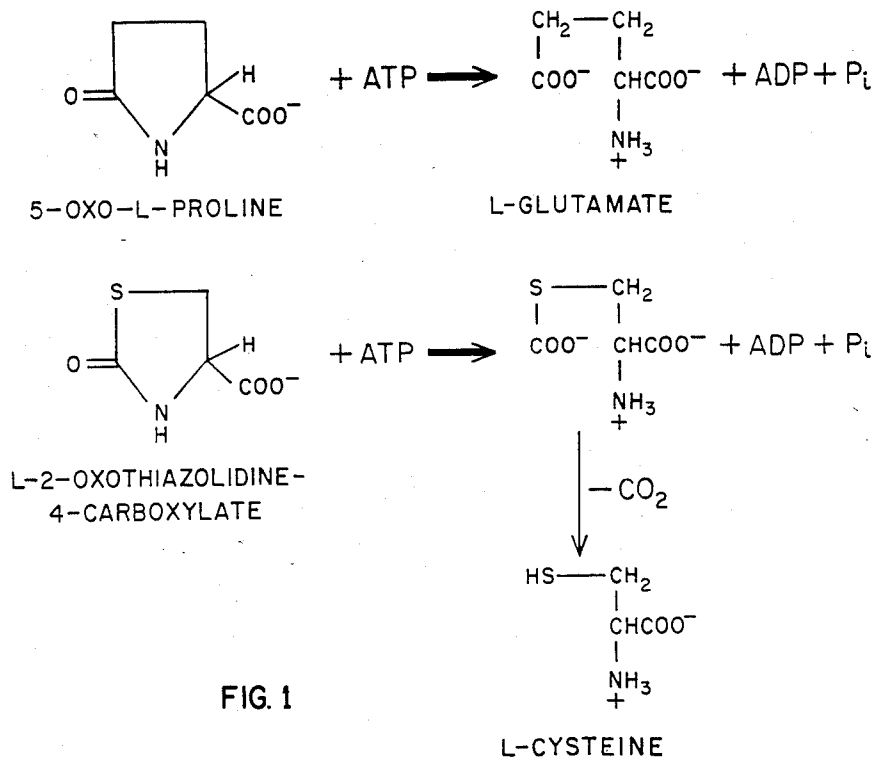


FIG. 1

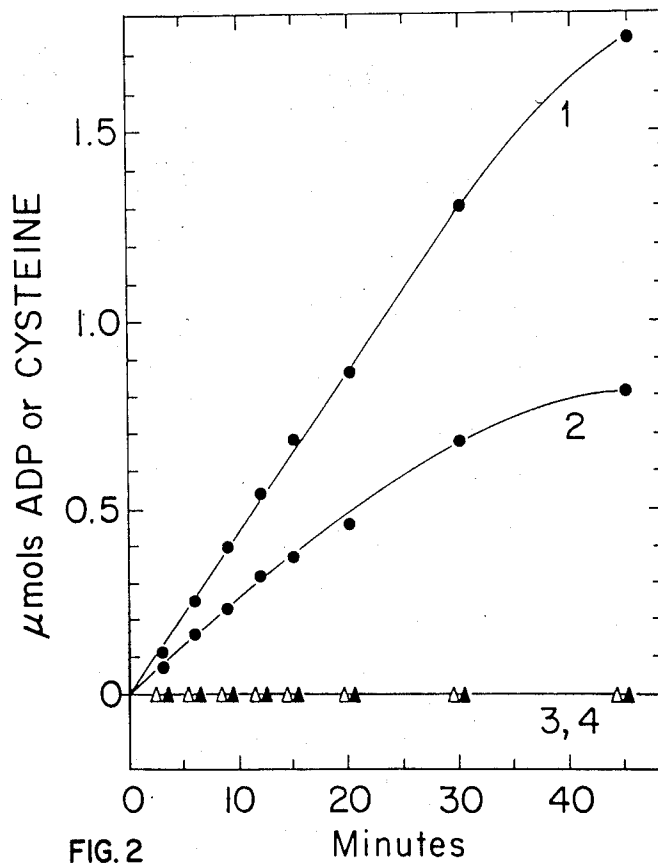
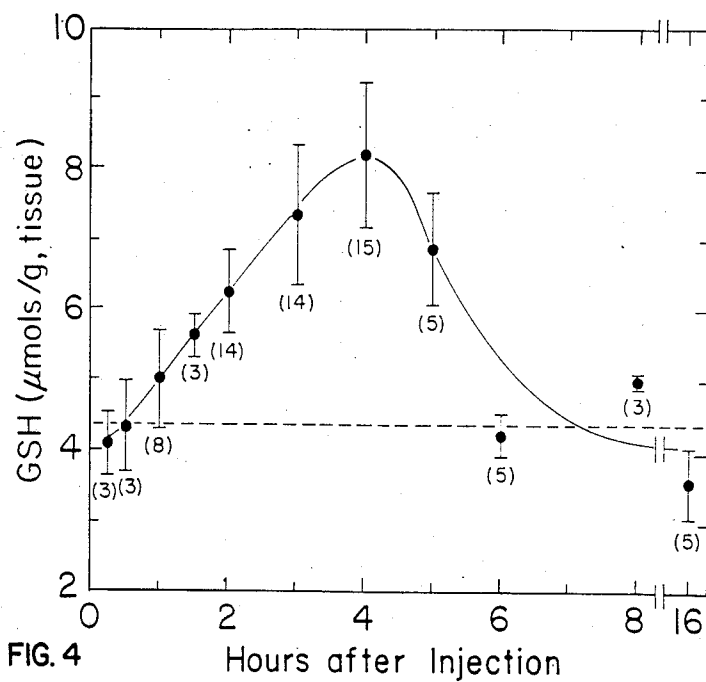
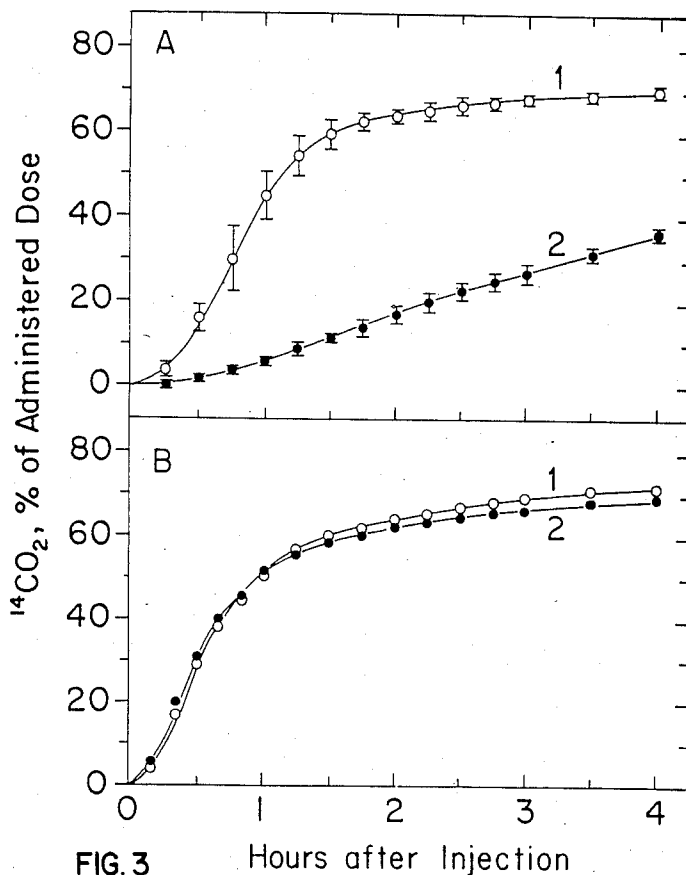


FIG. 2



CYSTEINE DELIVERY SYSTEM

The United States of American has rights to this invention pursuant to Public Health Grant AM-12034

This application is a continuation of application Ser. No. 547,459, filed Nov. 1, 1983, now abandoned.

This invention relates to a novel cysteine delivery system which is useful for introducing cysteine into *vivo* cells.

Specifically, this invention relates to a novel base amino acid solution (given orally or parenterally) and to a novel procedure for the production of cysteine within living cells.

The invention provides a method of restoring the glutathione level of numerous tissues where 5-oxoprolinase is present, particularly the human liver. The invention also provides a method of combating poisoning associated with the lessening or depletion of the glutathione content of cells. It is particularly applicable to the treatment of patients suffering from overdoses of acetaminophenols.

The nature of the invention will be more readily understood from the description to follow.

THE CHEMISTRY INVOLVED

It is known that 5-oxo-L-prolinase(L-pyroglutamate hydrolase) catalyzes the adenosine triphosphate (ATP)-dependent cleavage of 5-oxo-L-proline to glutamate. The products of such cleavage viz., glutamate, ADP (adenosine diphosphate) and P_i (inorganic ortho phosphate) are formed in a 1:1:1 ratio. It has now been found that a sulfur analog of 5-oxo-L-proline, viz, L-2-oxothiazolidine-4-carboxylate is also acted upon by this enzyme but the products include cysteine rather than glutamate. In the latter reaction the carboxylate is cleaved to yield an ADP/cysteine ratio of 1:1. The enzyme which exhibits an affinity for the analog similar to that for the natural substrate is inhibited by the analog, both *in vitro* and *vivo*. Thus, the carboxylate serves as a potent inhibitor of the gamma-glutamyl cycle at the step of 5-oxoprolinase. Administration of L-2-oxothiazolidine-4-carboxylate to mice that had been depleted of hepatic glutathione led to the restoration of normal hepatic glutathione levels. Since L-2-oxothiazolidine-4-carboxylate is an excellent substrate of the enzyme, it has been found to be useful as a component of base amino acid nutritional solutions and as an intracellular delivery system for cysteine. It is a good therapeutic agent for correcting conditions causing glutathione lessening or depletion, e.g., that resulting from poisoning induced by excessive amounts of acetaminophenol in the system.

The chemical reactions involved in the enzyme activity discussed above are shown in FIG. 1.

FIG. 2 is a chart showing the rate of formation of ADP and cysteine; and the absence of reaction when ATP was omitted or when the L-isomer was replaced with the D-isomer.

BASE AMINO ACID SOLUTIONS

Approximately 50% of hospitalized patients require some degree of nutritional support to counteract malnutrition and to assist in disease recovery. Many of such patients cannot take food directly because of G. I. tract dysfunction or poor oral intake. One of the primary means for establishing total parenteral nutrition is the infusion of basic amino acid solutions via central vein or

peripheral vein administration. Typical base amino acid solutions contain eight essential amino acids and seven non-essential amino acids and are normally prepared and sold commercially in 10% aqueous solutions. Of the amino acids not contained in such solutions the most important is cysteine. As will be explained below, cysteine cannot be administered intravenously due to its toxic effects on the system. Cysteine is quite important to human metabolism since, of all the amino acids contained in the basic solutions, the only one containing sulphur is L-methionine. Methionine is metabolized only in the liver so that if a patient has a malfunctioning or partially functioning liver, his system becomes devoid of sulphur bearing protein. The present invention provides a method for introducing cysteine which can be metabolized as well in organs other than the liver.

In accordance with one aspect of this invention the basic amino acid solutions are modified by the incorporation therein of 0.25 to 2.5 g/dl (grams per deciliter) of L-2-oxothiazolidine-4-carboxylate. A typical standard crystalline amino acid injection (10 wt.% solution) for central vein administration contains the following ingredients:

Essential Amino Acids	g/dl	Non-Essential Amino Acids	g/dl
L-Isoleucine	0.720	L-Alanine	1.280
L-Leucine	0.940	L-Arginine	0.980
L-Lysine (as acetate salt)	0.720	L-Histidine	0.300
L-Methionine	0.400	L-Proline	0.860
L-Phenylalanine	0.440	L-Serine	0.420
L-Threonine	0.520	L-Tyrosine	0.044
L-Tryptophan	0.160	Aminoacetic Acid (Glycine)	1.280
L-Valine	0.800		

Another standard base amino acid solution for central vein administration contains only the essential amino acids in 5.3 wt. % concentration.

A standard base amino acid formulation for peripheral vein injection is a 3.5 wt.% solution of the essential and non-essential amino acids and electrolytes, such as sodium, potassium, magnesium, chloride and acetate.

To the above nutrition support solutions there is added in accordance with this invention L-2-oxothiazolidine-4-carboxylate, preferably in the form of its neutral salt e.g., sodium, potassium, magnesium, etc., in the amount specified above.

EXPERIMENTS

In the experiments the reaction mixtures were made up to contain (final volume 0.5 ml) 100 mM Na Hepes (Na salt of N-2-hydroxyethylpiperazine-N'-2 ethanesulfonic acid) buffer (pH 8.0), 150 mM KCl, 8 mM MgCl₂, 2 mM phosphoenolpyruvate, 5 mM ATP (adenosine triphosphate), pyruvate kinase, in excess, 20 micrograms of 5-oxoprolinase, and the substrate as indicated.

The L- and D-isomers of 2-oxothiazolidine-4-carboxylate were synthesized by the method set out in Kaneko et al, Bull, Chem. Soc. (Japan) vol. 37, pp. 242-4 (1964) as modified by the method of Shah et al, Cancer Research vol. 39, pp. 3942-7 (1979). 5-Oxo-L-prolinase was isolated from rat kidney and freed of 5-oxoprolinase by gel filtration. Other chemicals and materials were obtained commercially.

After incubation for 30 min. at 37° C., the reaction mixtures were treated with 0.1 volume of 1M HCl, and placed at 0° C. for 5 min.; and an equivalent volume of 1M 2-amino-2-hydroxymethyl-1,3-propanediol was

added. Portions were analyzed for amino acids by use of a Durrum Model 500 amino acid analyzer. Cysteine was also determined in some experiments by reaction with 5,5'-dithiobis(2-nitrobenzoate) (DTNB).

RESULTS

When the enzyme 5-oxo-L-prolinase was incubated with L-2-oxothiazolidine-4-carboxylate in the presence of ATP there was rapid formation of ADP and cysteine. Energy for the reaction is supplied by the cleavage of ATP. No reaction was observed when ATP was omitted or when the L-isomer was replaced by the corresponding D-isomer. The ratio of ADP to cysteine produced increased with time due undoubtedly to the loss of cysteine due to oxidation. A more quantitative characterization of the reaction was obtained in studies in which the formation of cysteine was estimated by determining derivatives of this amino acid product, wherein the formation of ADP and cysteine (determined, after oxidation, as cystine, or after reduction and derivatization as the corresponding S-acetamido compound) was found to be stoichiometric. It was noted that the apparent K_m value for the L-isomer is somewhat lower than that found for the natural substrate, 5-oxo-L-proline; however, the calculated V_{max} value for the natural substrate is higher than that found for the L-isomer. As expected, L-2-oxo-thiazolidine-4-carboxylate is an excellent inhibitor of the utilization of 5-oxo-L-proline by the purified enzyme; thus, in reaction mixtures containing 1 mM 5-oxo-L-proline, the addition of L-2-oxothiazolidine-4-carboxylate at concentration of 1, 5, and 10 mM produced 73, 92 and 97% inhibition of glutamate formation, respectively. Under these conditions D-2-oxothiazolidine-4-carboxylate did not inhibit glutamate formation.

Utilizing reactions mixtures comparable to that described above tests were made on the natural substrate and on three substrate analogs. The results are summarized in the following table.

TABLE 1

Substrate (conc.,mM)	Action of 5-Oxo-L-Prolinase on Substrates	
	Products Formed ADP ^(a)	(nMoles) Amino Acid
5-Oxo-L-proline (2)	924	916 ^(b)
L-2-oxothiazolidine-4-carboxylate (2)	478	466 ^(c)
D-2-oxothiazolidine-4-carboxylate (5)	499	463 ^(d)
L-2-imidazolidone-4-carboxylate (2)	96	0 ^(e)
L-2-imidazolidone-4-carboxylate (2)	930	0 ^(f)

^(a)Determined by pyruvate kinase-lactate dehydrogenase coupled assay.

^(b)as glutamate;

^(c)determined after reduction with KBH_4 and reaction with iodoacetamide;

^(d)after conversion to cystine;

^(e)less than 20 nmol;

^(f)less than 5 nmol.

FIG. 2 shows graphically the results obtained with reaction mixtures similar to those of Table 1 except as follows: In Curves 1 and 2, 5 mM of the L-isomer and 5 mM ATP were contained; in Curve 3, 5 mM of the D-isomer and 5 mM ATP in Curve 4, 5 mM of the L-isomer but no ATP was added. At the indicated intervals portions were analyzed for ADP by the method indicated in the Table footnote, and for cysteine by reaction with DTNB.

The in vitro results showing that L-2-oxothiazolidine-4-carboxylate inhibits utilization of 5-oxoproline were confirmed by in vivo experiments. In the latter, animals were injected with 5-oxo-L-[^{14}C]proline and the respi-

ratory [$^{14}CO_2$] produced was collected and measured for several hours. The results show that a marked decrease in the formation of [$^{14}CO_2$] occurred in the animals treated with L-isomer which excreted about 16% of the injected dose in their urine; no radioactivity was found in the urine of the controls. Upon injection of L-[^{14}C]glutamate to the animals the findings indicate that injection of L-2-oxothiazolidine-4-carboxylate does not have a significant effect on the metabolism of glutamate. Previous studies of the inventors showed that injection of another analog viz., L-2-imidazolidone-4-carboxylate also markedly inhibits the in vivo metabolism of 5-oxoproline, but not that of glutamate. However, L-2-oxothiazolidine-4-carboxylate is a much more efficient inhibitor. Comparison of the effects of the thiazolidine compound with those of L-2-imidazolidone-4-carboxylate indicate that the thiazolidine compound is at least four times more active on a molar basis.

ANIMAL STUDIES

The effect of injecting mice with L-2-oxothiazolidine-4-carboxylate on the level of hepatic glutathione was studied. The findings indicate that administration of the thiazolidine stimulated the formation of glutathione to levels that are about twice those of the controls. The maximal effect was observed about 4 hours after injection. No such stimulation was observed when the animals were injected with equivalent doses of D-2-oxothiazolidine-4-carboxylate. These findings are indicative of in vivo formation of cysteine after administration of L-2-oxothiazolidine-4-carboxylate. No free cysteine was detected in acid extracts of the livers of control or experimental animals by amino acid analyses carried out after reduction by dithiothreitol and derivatization with 2-vinylpyridine. This finding, not surprising in view of the very low levels of tissue cysteine, indicates that cysteine formed by cleavage of the thiazolidine is rapidly utilized for glutathione syntheses.

FIG. 3 depicts the inhibition of 5-oxoproline metabolism by L-2-oxothiazolidine-4-carboxylate. Mice fasted overnight were injected intraperitoneally with NaCl (Curve 1; open symbols; controls) or with Na L-2-oxothiazolidine-4-carboxylate (2.5 mmol/kilo) (Curve 2; closed symbols). After 10 min., the mice were injected subcutaneously with 5-oxo-L-[^{14}C]proline (A) or with L-[^{14}C]glutamate (B) (0.02 mmol; 1.1×10^6 cpm). The mice were then placed in metabolic chambers and the respiratory CO_2 was collected in ethanolamine/methanol, 1:4 (vol/vol). In A, each curve represents the results on 3 animals; the points are mean values and the bars represent standard deviations from the mean. In B, each curve represents the average of closely-agreeing results obtained from 2 animals.

FIG. 4 shows the increase of hepatic levels of glutathione after injection of L-2-oxothiazolidine-4-carboxylate. Mice fastened overnight were injected intraperitoneally with NaCl (controls; dashed line) or with Na L-2-oxothiazolidine-4-carboxylate (6.5 mmol/kilo). At intervals, the mice were sacrificed and the excised liver was homogenized in 5 volumes of 1% picric acid. After centrifugation, the protein-free supernatant solutions were analyzed for glutathione by the method of Tietze, Anal. Biochem 27, 502-22 (1969). Analyses were carried out in duplicate. In a number of instances, these values were checked by the 2-vinylpyridine procedure of Griffith, Anal. Biochem. 106; 207-12 (1980); good agreement between the two methods was observed. In the figure, the points are mean values and the bars indi-

cate the average deviation from the mean. The numbers in parentheses give the number of animals used. The value for the controls was 4.40 ± 0.9 $\mu\text{mol/g}$.

Administration of L-2-oxothiazolidine-4-carboxylate to animals leads to an increase in the cysteine content of tissues, especially the liver. Administration of this compound therefore provides a way of getting cysteine into cells. One cannot give cysteine itself without complications since this amino acid is toxic extracellularly. However, when the L-oxothiazolidine compound is injected it enters the cell where it is split giving rise to cysteine within the cell. In a number of clinical situations, for example, after poisoning with excessive pain-killing drugs, e.g., those containing active ingredients known as N-acetyl-para-aminophenol, paracetamol, and acetaminophen, the liver glutathione levels decrease markedly. When cysteine is supplied to liver cells, additional glutathione can be synthesized and thus serve usefully in detoxifying the offending drug. Therefore, L-2-oxothiazolidine-4-carboxylate serves as an intracellular delivery system for cysteine and has potential as a therapeutic agent in situations in which there is depletion of liver glutathione. After acetoaminophenol is given to experimental animals such as mice, the liver glutathione declines rapidly because it is extensively utilized by reaction with the drug. If one gives the L-isomer compound even two hours after the drug is given, an increase in liver glutathione occurs indicating the potential value of this compound in treatment. For human treatment the L-isomer, preferably in the form of its neutral salt is administered directly into the gastro-intestinal tract or introduced intravenously.

Having described the invention so that it may be practiced by those skilled in the art: what is claimed is:

1. A method for introducing cysteine into in vivo cells containing 5-oxo-L-prolinase which comprises introducing into said cells L-2-oxothiazolidine-4-carboxylate in an amount sufficient to be converted in the presence of 5-oxo-L-prolinase to cysteine, and thereby form increased levels of cysteine in said cells.

2. A method for introducing cysteine into mammalian cells containing 5-oxo-L-prolinase which comprises orally or parenterally administering to a mammal containing said cells L-2-oxothiazolidine-4-carboxylate in an amount sufficient to be converted in the presence of 5-oxo-L-prolinase to cysteine, and thereby form increased levels of cysteine in said cells.

3. A method according to claim 2 in which said mammal is a human.

4. A method for introducing cysteine into mammalian cells containing 5-oxo-L-prolinase which comprises parenterally administering to a mammal containing said cells L-2-oxothiazolidine-4-carboxylate in an amount sufficient to be converted in the presence of 5-oxo-L-prolinase to cysteine, and thereby form increased levels of cysteine in said cells.

5. A method according to claim 4 in which said mammal is a human.

6. A method according to claim 4 wherein said L-2-oxothiazolidine-4-carboxylate is administered intravenously to a mammal.

7. A method according to claim 6 wherein said L-2-oxothiazolidine-4-carboxylate is administered in form of an aqueous solution containing from 0.25 to 2.5 grams per deciliter of L-2-oxothiazolidine-4-carboxylate.

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UNITED STATES PATENT AND TRADEMARK OFFICE
CERTIFICATE OF CORRECTION

PATENT NO. : 4,665,082

DATED : May 12, 1987

INVENTOR(S) : Meister et al.

It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

Title page, item [63], under "Related U.S. Application Data", after "abandoned", please delete the period and insert ", which is a continuation of application Serial No. 388,907, filed April 15, 1982, abandoned, which is a division of application Serial No. 233,564, filed February 11, 1981, Patent No. 4,335,210."

Signed and Sealed this

Twenty-fourth Day of December, 1996

Attest:



BRUCE LEHMAN

Attesting Officer

Commissioner of Patents and Trademarks